GREENEX ANTIBACTERIAL- chloroxylenol liquid
Cleanslate Group LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Greenex Foam Hand Soap

Drug Facts

Active Ingredient

Parachlorometaxylenol 0.3% w/w

Purpose

Antiseptic

Uses

- Handwash to help reduce bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

Warnings

- **For external use only.**
- Keep out of eyes, ears or mouth. In case of eye contact, flush eyes with water.
- **Stop use and ask a doctor** if irritation or redness develop or if condition persists for more than 72 hours.
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands with water and dispense sufficient amount of product into cupped palm of hand.
- Wash both hands thoroughly for 15 seconds.
- Rinse under running water and dry thoroughly.

Inactive Ingredients

Citric acid, Cocamide DEA, DMDM Hydantoin, Ethyl Alcohol, FD&C Red 4, Fragrance, Isopropyl Alcohol, Phenoxyethanol, Sodium Laureth Sulfate, Water.

GREENEX

2 Bergen Turnpike

Ridgefield Park, NJ 07660

MADE IN THE USA

GREENEX

ANTIBACTERIAL FOAM HAND SOAP

With 0.3% PCMX

1000 ml

GREENEX ANTIBACTERIAL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80586-513
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALCOHOL (UNII: 3K9958V90M)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80586-513-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/13/2021	

Labeler - Cleanslate Group LLC (117657934)

Revised: 4/2021

Cleanslate Group LLC